



## Environmental controls of greenhouse gas release in a restoring peat bog in NW Germany

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# Environmental controls of greenhouse gas release in a restoring peat bog in NW Germany

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## Abstract

In Central Europe, most bogs have a history of drainage and many of them are currently being restored. Success of restoration as well as greenhouse gas exchange of these bogs is influenced by environmental stress factors as drought and atmospheric nitrogen deposition. We determined the methane and nitrous oxide exchange of sites in the strongly decomposed center and less decomposed edge of the Pietzmoor bog in NW Germany in 2004. Also, we examined the methane and nitrous oxide exchange of mesocosms from the center and edge before, during, and following a drainage experiment as well as carbon dioxide release from disturbed unfertilized and nitrogen fertilized surface peat. In the field, methane fluxes ranged from 0 to  $3.8 \text{ mg m}^{-2} \text{ h}^{-1}$  and were highest from hollows. Field nitrous oxide fluxes ranged from 0 to  $574 \mu\text{g m}^{-2} \text{ h}^{-1}$  and were elevated at the edge. A large *Eriophorum vaginatum* tussock showed decreasing nitrous oxide release as the season progressed. Drainage of mesocosms decreased methane release to 0, even during rewetting. There was a tendency for a decrease of nitrous oxide release during drainage and for an increase in nitrous oxide release during rewetting. Nitrogen fertilization did not increase decomposition of surface peat. Our examinations suggest a competition between vascular vegetation and denitrifiers for excess nitrogen. We also provide evidence that the von Post humification index can be used to explain greenhouse gas release from bogs, if the role of vascular vegetation is also considered. An assessment of the greenhouse gas release from nitrogen saturated restoring bogs needs to take into account elevated release from fresh *Sphagnum* peat as well as from sedges growing on decomposed peat. Given the high atmospheric nitrogen deposition, restoration will not be able to achieve an oligotrophic ecosystem in the short term.

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## 1 Introduction

Due to the high amount of carbon stored in the peatlands of the world and the sensitivity of biogeochemical processes in these ecosystems to climate change, research on matter cycling in peatlands has received considerable interest. Especially the release of greenhouse gases (GHG) as carbon dioxide (CO<sub>2</sub>), methane (CH<sub>4</sub>), and nitrous oxide (N<sub>2</sub>O) from peatlands has been the focus of biogeochemical research due to its potential contribution to feedbacks to global warming. Despite large areas of (often degraded) peat bodies in temperate regions, research on peat bogs is mostly from natural boreal sites (Blodau, 2002).

In temperate Germany, widespread drainage of bogs resulted in a serious decline of peatland area. Today, in NW Germany (Lower Saxony) merely 5% of formerly 2348 km<sup>2</sup> bog area remain undisturbed or in a close to natural state (Schmatzler, 1990). Therefore, protection of the remaining intact peat bogs is accompanied by restoration efforts in moderately degraded bogs. The most important environmental constraints on the successful restoration of these bogs are i) a low water table, a result of previous drainage and climate change, ii) atmospheric N deposition, and iii) strong decomposition of degraded peat.

The importance of water table on CO<sub>2</sub>, CH<sub>4</sub>, and N<sub>2</sub>O release from peat has been discussed extensively (Blodau, 2002). It has been reported that CO<sub>2</sub> evolution follows an optimum function, with highest rates at an intermediate water table (Glatzel et al., 2006). Magnitude and important parameters of CH<sub>4</sub> emission from wetlands are well known (Le Mer and Roger, 2001). Drainage decreases CH<sub>4</sub> release and rewetting does not necessarily lead to an immediate rise in CH<sub>4</sub> release (Tuitilla et al., 2000).

Nutrients that may limit decomposition include nitrogen (N) and phosphorus (P) (Güsewell and Freeman, 2003). In Lower Saxony, even “undisturbed” bogs are subject to elevated N deposition of up to 70 kg ha<sup>-1</sup> yr<sup>-1</sup> (Gauger et al., 2002). At these high N deposition rates, the capacity of *Sphagnum* to take up N is exceeded (Lamers et al., 2000), N concentration in pore water accumulates and plants with high N demand

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as *Molinia caerulea* become more competitive (Limpens et al., 2003). An increasing proportion of easily decomposable litter, and N enriched *Sphagnum* tissue enhance decomposition and N mineralization (Lamers et al., 2000; Aerts et al., 1992), facilitating N<sub>2</sub>O and CO<sub>2</sub> release. Generally, in bogs, N<sub>2</sub>O release is most common in disturbed locations influenced by elevated N content (Regina et al., 1996). N<sub>2</sub>O production requires the availability of nitrogen and is highest at high soil moisture, but not inundation (Granli and Bøckmann, 1994). A direct influence of atmospheric NO<sub>3</sub> deposition on N<sub>2</sub>O release has been reported by Aerts (1997) and Hefting et al. (2003).

Decomposition status of peat controls its potential for further decomposition. Examinations by Glatzel et al. (2004) demonstrated a decreasing potential for aerobic and anaerobic CO<sub>2</sub> and CH<sub>4</sub> production with a rising von Post decomposition index. Moore and Dalva (1997) were not able to relate aerobic CO<sub>2</sub> production to the degree of decomposition of 140 peat samples. In the Pietzmoor Glatzel et al. (2006) explained increased CO<sub>2</sub> release from *Sphagnum* hollow peat compared to hummock peat by lower decomposition rates of hollow peat. Alm et al. (1999) remarked that increased NO<sub>3</sub> availability may be due to high decomposition, increasing rates of N<sub>2</sub>O emission from drained peatlands.

In this contribution we intend to clarify the influence of these controls on the GHG release of a restoring temperate bog. Previous investigations (Glatzel et al., 2006) have shown the effect of drought on decomposition rates. Specifically, we investigate the influence of a drawdown in water table and peat properties on methane and nitrous oxide release in a restoring peat bog and the influence of nitrogen on decomposition of surface peat. We hypothesize that i) drought decreases the CH<sub>4</sub> and N<sub>2</sub>O release in the bog and rewetting temporarily increases CH<sub>4</sub> and N<sub>2</sub>O release, ii) decomposition of peat controls CH<sub>4</sub> and N<sub>2</sub>O release, and iii) atmospheric nitrogen deposition accelerates decomposition of surface peat.

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## 2 Site and methods

### 2.1 Research site

The study site was the Pietzmoor (Lower Saxony; NW, Germany; 53°06' N; 9°50' E). The bog is located on the eastern edge of the closed occurrence of rainfed bogs in NW Germany. Mean annual precipitation is 790 mm; mean annual temperature is 8°C. The examination period was March to September 2004. Atmospheric N deposition is ca. 22 kg ha<sup>-1</sup> yr<sup>-1</sup> (Fottner et al, 2004). Today, the Pietzmoor is moderately degraded. Manual peat extraction at the edges of the Pietzmoor was conducted between the 16th century and 1960. Deep drainage ditches, constructed in the 19th century further degraded the bog, resulting in increased growth of birch (*Betula* sp.) and pine (*Pinus* sp.). Since 1970, when restoration efforts began, drainage ditches have been closed and trees cut. This resulted in formation of a recent superficial acrotelm with *Sphagnum* spp. growing in many hollows. Hummocks are still dominated by *Empetrum nigrum*, *Calluna vulgaris*, and *Eriophorum vaginatum*.

### 2.2 Field CH<sub>4</sub> and N<sub>2</sub>O flux determination

Between March and August 2004, CH<sub>4</sub> and N<sub>2</sub>O fluxes were determined 14 times employing a closed chamber method (Hutchinson and Livingston, 1993) at 10 locations within the Pietzmoor bog. Of the 10 previously installed collars (covering 0.068 m<sup>-2</sup>), five collars were installed in the center and at the edge of the bog. Six collars covered hummocks and four collars covered hollows. Among the hollow collars, two were vegetated by *Sphagnum fallax*, one hosted a small *Eriophorum vaginatum* tussock and one contained no living vegetation. Among the hummock collars, three were vegetated by *Calluna vulgaris*, one contained a big and one a small *Eriophorum vaginatum* individuum, and one was inhabited by lichens. These collars covered the range of microsites in the bog previously determined by Rathert (2004).

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For gas flux determination, gas samples from the closed chamber were sampled by syringe five times in 5 min intervals and transported to the laboratory in Göttingen. The syringes were attached to an autosampler coupled to a Shimadzu GC-14B gas chromatograph and a set of four different calibration gas cocktails (described by Lottfield et al., 1997). Precision of analysis was 0.4% for CH<sub>4</sub> and 1.0% for N<sub>2</sub>O. As no saturation effects were found, fluxes were calculated from the linear slope of the concentration change over time (Lessard et al., 1994).

### 2.3 CH<sub>4</sub> and N<sub>2</sub>O release from mesocosms

Twelve undisturbed peat cores (diameter 15 cm) were sampled by cutting the peat at the outside of tube and simultaneously pushing the tube above the cut peat until average 23 cm of peat were inside the tube. All cores were taken from hollows, six in the centre and six from the edge. The peat cores were transferred into 30 cm high mesocosms that enabled sampling of percolating water and gas concentrations from a 7 cm headspace. Peat cores were watered in three day intervals with artificial Schneverdingen rain (diluted ammonium nitrate solution set to a pH of 4.5, equivalent to an amount of 790 mm yr<sup>-1</sup> and 20 kg dry and wet N deposition ha<sup>-1</sup> yr<sup>-1</sup>). As suggested by Blo-dau et al. (2004), a two month equilibration phase preceded the experiment. During the equilibration phase, the water table was set to 7 cm below ground. The cores were stored at 20°C close to windows, allowing a natural night and day regime. Vegetation (*Sphagna* and small herbs, no large plants) continued to grow during the experiment.

The experiment consisted of three phases. The pre-drainage phase preceded the drainage phase. During this phase, the six manipulated cores were subjected to free drainage (restricted to 100 mL d<sup>-1</sup>) without applying low pressure. At the control cores water table remained close to the peat surface. During the second phase (drainage phase), the manipulated cores were subjected to free drainage. The third phase (post-drainage phase) began by closing the drainage at experimental cores and the daily addition of 40 mL artificial Schneverdingen rain until the water table was back to 7 cm below ground. The pre-drainage phase lasted 5 to 8 days, the drainage phase until

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the elimination of standing water lasted 5 to 6 days and the regeneration of high water table (post drainage phase) took 12 to 14 days.

During the experiment, we determined gas fluxes from all cores as described above (except for a 30 s sampling interval due to the small headspace) daily. Following the experiment, carbon (C) and N concentration of peat from all cores was determined. This was done by drying peat at 45°C from all horizons, milling it to 0.25 mm and analysis by combustion at 900°C in a LECO CN- Analyzer (LECO, St. Joseph, MI, USA). The C and N concentration of all horizons were averaged to 0–15 cm depth. We also estimated the von Post humification index at all cores.

## 2.4 CO<sub>2</sub> evolution from incubated disturbed samples

We sampled peat from 0–10 cm depth from *Calluna* hummocks and *Sphagnum* hollows in the Pietzmoor. Approximately 20 g of peat were set to 75% water content, which yields intermediate rates of CO<sub>2</sub> evolution (Glatzel et al., 2006) and placed in 400 mL jars in triplicate. All samples were additionally moistened by 1 mL of liquid. The fertilized samples received 0.036 M ammonium nitrate solution (equivalent to 50 kg N ha<sup>-1</sup>), and the unfertilized control samples received plain water.

The incubation experiment was conducted using the method by Isermeyer (1952) following the experimental design described Glatzel et al. (2006). Briefly, evolved CO<sub>2</sub> was absorbed by 20 mL of 0.1 M NaOH adsorption inside the jars. Sampling of NaOH placed in small containers) following 1, 3, 6, 11, 17, 28 and 42 of incubation and titration with 0.1 M HCl allows the calculation of CO<sub>2</sub> evolved since the preceding sampling date.

## 2.5 Ancillary measurements and statistical procedures

We measured air temperature and precipitation at a weather station located 2 km from the field site and installed an air temperature logger 20 cm above the surface of the bog. We determined water table by previously installed wells 14 times between March and August 2004 at the center and the edge of the Pietzmoor. All data sets were tested for

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normal distribution using the Kolmogorov-Smirnov test. Data on N<sub>2</sub>O release and day of year (Fig. 3) was normal distributed, so Pearson's correlation coefficient was calculated. The other data was generally not normal distributed, and  $n$  was generally small, so correlation analyses were carried out using Spearman's rho test and differences between data subsets were analyzed using the Wilcoxon test employing the Statistica 6.1 software package (Stat Soft, 2004).

### 3 Results

#### 3.1 Weather and water table

The field season was warmer and wetter than the long term mean (1989 to 2004). Between March and August 2004, we recorded 427 mm precipitation as opposed to a long term mean of 381 mm. Mean temperature during the field season was 14.2°C, compared a long term mean of 13.8°C. At the start of the field season, water table was close to the surface (Fig. 1). Following a rather dry spring, frequent precipitation led to a rise in water table until early July. In July and August, water table dropped to 25 cm, but rose again in late August. In the center of the bog, water table responded more quickly than at the edge.

#### 3.2 Field CH<sub>4</sub> and N<sub>2</sub>O fluxes

Field CH<sub>4</sub> fluxes ranged from 0 to 7.8 mg CH<sub>4</sub> m<sup>-2</sup> h<sup>-1</sup> and averaged 1.2 mg CH<sub>4</sub> m<sup>-2</sup> h<sup>-1</sup>. Spatial variability of CH<sub>4</sub> fluxes was high, so we were not able to detect significant differences between the mean CH<sub>4</sub> flux from hummocks and hollows and between the collars in the center and at the edge of the bog (Fig. 2), although there was a tendency for elevated CH<sub>4</sub> release in hollows and at the center of the bog. As the water table at the center was not lower than at the edge, the absence of a significant difference between CH<sub>4</sub> release at the two sites is not surprising.

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Although N<sub>2</sub>O fluxes in the field were generally low, and often 0 at some collars, we detected a N<sub>2</sub>O release of up to 574 μg m<sup>-2</sup> h<sup>-1</sup>. We found no N<sub>2</sub>O uptake. There was no difference in N<sub>2</sub>O release between hummocks and hollows, but at the edge, nitrous oxide release was higher than at the center despite the lack of a difference in water table (Fig. 1).

During the course of the season, CH<sub>4</sub> fluxes rose from 0.5 mg m<sup>-2</sup> h<sup>-1</sup> to 2 mg m<sup>-2</sup> h<sup>-1</sup> (at some hummocks) and 4 to 8 g m<sup>-2</sup> h<sup>-1</sup> (at some hollows). This trend could not be noticed at all collars. There was no seasonal trend of N<sub>2</sub>O fluxes, except for the collar vegetated by a large *Eriophorum vaginatum* tussock. There, N<sub>2</sub>O fluxes decreased linearly with the course of the season (Fig. 3).

### 3.3 CH<sub>4</sub> and N<sub>2</sub>O release from mesocosms

#### 3.3.1 Methane

Methane release from the cores was higher than from field sites, averaging 8.2 mg CH<sub>4</sub> m<sup>-2</sup> h<sup>-1</sup>. During the pre-drainage phase, there was no difference in CH<sub>4</sub> flux between the control cores and the manipulated cores. During this phase, methane fluxes were between 0.1 and 84.5 mg m<sup>-2</sup> h<sup>-1</sup> and averaged 7.6±9.1 to 8.7±11.7 mg CH<sub>4</sub> m<sup>-2</sup> h<sup>-1</sup> (Fig. 4). During drainage, the control cores remained at the CH<sub>4</sub> release level, emitting -0.1 to 138 mg CH<sub>4</sub> m<sup>-2</sup> h<sup>-1</sup> and averaging 9.4±11.7 mg CH<sub>4</sub> m<sup>-2</sup> h<sup>-1</sup>. Methane release of the manipulated cores dropped to 0 to 3.1 mg CH<sub>4</sub> m<sup>-2</sup> h<sup>-1</sup> with a mean release of 0.3±0.4 mg CH<sub>4</sub> m<sup>-2</sup> h<sup>-1</sup>. During the third phase, CH<sub>4</sub> emissions from the control plots remained at 0 to 99.6 mg CH<sub>4</sub> m<sup>-2</sup> h<sup>-1</sup> an average value of 8.3±12.9 mg CH<sub>4</sub> m<sup>-2</sup> h<sup>-1</sup>. During the post-drainage phase, CH<sub>4</sub> emissions from the manipulated cores remained at the level of the drainage phase emitting 0 to 11.2 mg CH<sub>4</sub> m<sup>-2</sup> h<sup>-1</sup> an a mean CH<sub>4</sub> release of 0.3±0.3 mg m<sup>-2</sup> h<sup>-1</sup>. In summary, CH<sub>4</sub> release of the manipulated cores remained at close to zero even when the water table reached the original position.

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### 3.3.2 Nitrous oxide

Variability of emissions of  $\text{N}_2\text{O}$  from the cores was higher than the variability of  $\text{CH}_4$  emissions. During the first phase,  $\text{N}_2\text{O}$  release from the control cores was 0 to  $1571 \mu\text{g m}^{-2} \text{h}^{-1}$  (Fig. 5). Previous to drainage, the manipulated cores released 0 to  $2255 \mu\text{g N}_2\text{O m}^{-2} \text{h}^{-1}$ . Thus, control cores released  $292 \pm 361 \mu\text{g N}_2\text{O m}^{-2} \text{h}^{-1}$  opposed to  $163 \pm 190 \mu\text{g N}_2\text{O m}^{-2} \text{h}^{-1}$  from the manipulated cores, possibly showing an (non significant) effect of beginning drainage. During the drainage phase,  $\text{N}_2\text{O}$  release at manipulated as well as control cores was lower than during the first phase. Due to the higher emission at the manipulated cores during the pre-drainage phase, this change was significant for the manipulated cores in contrast to the control. During this phase, control cores released 0 to  $673 \mu\text{g N}_2\text{O m}^{-2} \text{h}^{-1}$  and manipulated cores emitted 0 to  $348 \mu\text{g N}_2\text{O m}^{-2} \text{h}^{-1}$ . The average values were  $75 \pm 59$  and  $73 \pm 102 \mu\text{g N}_2\text{O m}^{-2} \text{h}^{-1}$  for the control and the manipulated cores, respectively. During the post-drainage phase,  $\text{N}_2\text{O}$  release from the control cores remained at 0 to  $1464 \mu\text{g N}_2\text{O m}^{-2} \text{h}^{-1}$ , with an average of  $72 \pm 69 \mu\text{g N}_2\text{O m}^{-2} \text{h}^{-1}$ . The manipulated cores emitted 0 to  $1590 \mu\text{g N}_2\text{O m}^{-2} \text{h}^{-1}$ , with a mean  $\text{N}_2\text{O}$  release of  $150 \pm 157 \mu\text{g m}^{-2} \text{h}^{-1}$ , showing a (non significant) sign of increasing  $\text{N}_2\text{O}$  release. In summary, the extremely high variability and the multiple controls of  $\text{N}_2\text{O}$  release lead to an incoherent emission pattern.

Thus, the mesocosm experiment produced a clear result for  $\text{CH}_4$  and no clear result for  $\text{N}_2\text{O}$ . A lasting suppression of  $\text{CH}_4$  release during a following drainage is not mirrored by a similar effect for  $\text{N}_2\text{O}$ , although there is a tendency for decreased  $\text{N}_2\text{O}$  release during drainage and possibly a somewhat increased  $\text{N}_2\text{O}$  release following drainage.

### 3.3.3 Properties of the peat cores

Simple measures of surface peat point towards stronger decomposition of peat in the center of the bog (Table 1): C and N content in the top 15 cm of the peat cores from

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the center of the bog were significantly higher than from the edge of the bog. There was no significant difference in the C/N ratio from the cores sampled at the center to the ones sampled at the edge of the bog, but cores from the edge tended towards a higher C/N ratio. As evidenced by the von Post index, peat from the bog center was more humified than peat at the bog edge.

### 3.4 CO<sub>2</sub> evolution from incubated disturbed samples

According to the incubation experiment, N fertilization of surface peat does not control potential CO<sub>2</sub> release. In contrast to sampling depth or peat type (*Calluna* hummock or *Sphagnum* hollow), a wide range of unfertilized and fertilized samples did not differ in the amount of CO<sub>2</sub> release. Following 42 days of incubation, unfertilized peat released 43.7±40.1 mg CO<sub>2</sub> per g of dry peat and fertilized peat released 43.0±45.9 mg CO<sub>2</sub> per g of dry peat (Fig. 6).

## 4 Discussion

### 4.1 Field CH<sub>4</sub> and N<sub>2</sub>O fluxes

The CH<sub>4</sub> fluxes that we measured in the Pietzmoor are within the range previously reported by many authors and recently reviewed by Le Mer and Roger (2001) and Whalen (2005). Although the high spatial variability of CH<sub>4</sub> fluxes impedes the interpretation of data, we discuss patterns of CH<sub>4</sub> release. The elevated CH<sub>4</sub> emissions from hollows at our sites are probably due to the proximity to the water table and a shallower aerobic zone of CH<sub>4</sub> oxidation (Pelletier et al., 2007; Strack et al., 2004). Furthermore, some of the hollows are covered with *Eriophorum vaginatum*. Vascular plants, especially sedges are known for high CH<sub>4</sub> release (Joabsson et al., 1999; Strack et al., 2006) and *Eriophorum vaginatum* tussocks are CH<sub>4</sub> emission hotspots as they provide substrate for methanogenesis and provide a pathway for CH<sub>4</sub> release (Tuitilla et al., 2000; Marinier et al., 2004). The somewhat elevated CH<sub>4</sub> emissions at

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the center of the bog cannot be explained by water table. However, due to the higher decomposition, field moisture could be higher in the center than at the edge. Only recently, Basiliko et al. (2007) state that mining, alteration and restoration modify the factors controlling CH<sub>4</sub> production, e.g. indicated by a strong influence of soil moisture content on CH<sub>4</sub> production at mined and restored sites while no such correlation could be found at natural sites. In contrast to the hot and dry summer of 2003, the wet summer of 2004 did not cause any drought stress and water table in the center of the bog remained at the same level as at the edge. There was no profound drawdown of the water table. So, water table did not control CH<sub>4</sub> release and the highest CH<sub>4</sub> release (7.8 mg m<sup>-2</sup> h<sup>-1</sup>) took place on 8 April 2004 with the water table at 24.5 cm below the surface. On the one hand, this is in contrast to the well established relationship between CH<sub>4</sub> release and water table (Moore and Knowles, 1989; Moore and Dalva, 1993). On the other hand, this high CH<sub>4</sub> release took place at just one occasion, from an *Eriophorum vaginatum* tussock located in a hollow.

We are not able to explain the (insignificantly) elevated CH<sub>4</sub> emission in the center of the bog. Following the reasoning of Glatzel et al. (2004), the low degree of humification of surface peat at the edge of the Pietzmoor as evidenced by the von Post index (Table 1) should favor elevated CH<sub>4</sub> emission at that subsite. Glatzel et al. (2004) presented a negative correlation between von Post index and anaerobic CH<sub>4</sub> production rate in surface peat from several locations in eastern Canada. Therefore, the degree of humification of surface peat does not control CH<sub>4</sub> release at the Pietzmoor or factors not determined by us need to be taken into account. Still, the determination of the von Post index is a fast field method to get information about the decomposition state of the peat, which is indirectly linked to peat carbon substrate quality. According to Guckland (2004), who used spectroscopic methods to determine aromatic compounds in DOC, these compounds are frequent in the top 10 cm of the Pietzmoor. Consequently, the peat carbon quality is not a good proxy for methanogenesis, which once more highlights the importance of *Eriophorum vaginatum* as supplier of easily degradable compounds (Saarnio et al. 2004).

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As oligotrophic peatlands are generally N limited, they are usually no sources of  $\text{N}_2\text{O}$  (Martikainen et al., 1993). Thus, the field  $\text{N}_2\text{O}$  fluxes reported in this contribution are high compared with these sites. However it must be taken into account that most studies from pristine oligotrophic peatlands are from boreal sites with rather low atmospheric N deposition (Nordin et al., 1998). Our site has a history of drainage, is located in the temperate zone, experiences high atmospheric N input and a rapid fluctuation in water table (Fig. 1), and, at drought conditions,  $\text{NO}_3\text{-N}$  concentrations of  $22 \pm 31 \text{ mg L}^{-1}$  (Glatzel et al., 2006). The  $\text{N}_2\text{O}$  release from the Pietzmoor is higher than the  $\text{N}_2\text{O}$  release from a restoring peat bog in S Germany, where Drösler (2005) determined an  $\text{N}_2\text{O}$  emission of 1 to  $31 \mu\text{g N}_2\text{O m}^{-1} \text{ h}^{-1}$ . According to our research, only cultivated or drained peatlands release  $>100 \mu\text{g N}_2\text{O m}^{-1} \text{ h}^{-1}$ . Regina et al. (1999) measured an  $\text{N}_2\text{O}$  release of  $440 \mu\text{g N}_2\text{O m}^{-1} \text{ h}^{-1}$  in a Finnish drained tall sedge fen. On the other hand, the same authors found that rewetting reduces  $\text{N}_2\text{O}$  release from a previously drained birch-pine fen from 50 to  $100 \mu\text{g m}^{-1} \text{ h}^{-1}$ . Cultivated sites on organic soils from NW Finland released 70 to  $170 \mu\text{g N}_2\text{O m}^{-1} \text{ h}^{-1}$  (Maljanen et al., 2001).

Water table also did not control  $\text{N}_2\text{O}$  release. This is not surprising as the field campaign was rather short and  $\text{N}_2\text{O}$  release is at its maximum in unsaturated soil (Granli and Bøckmann, 1994). Due to the infamously high spatial and temporal variability of soil  $\text{N}_2\text{O}$  emissions (Folorunso and Rolston, 1984), the lack of a difference of  $\text{N}_2\text{O}$  release between hummocks and hollows is not surprising. However, the significantly elevated  $\text{N}_2\text{O}$  release from the edge of the Pietzmoor compared to its center (Fig. 2) is surprising. Even when there is no difference in water table, nitrous oxide flux from the edge of the bog is elevated. Unfortunately, no N data from peat pore water are available from 2004. However, an increased peat pore water  $\text{NO}_3$  concentration at the edge of the bog is unlikely: In contrast to the center of the Pietzmoor,  $\text{NO}_3$  concentrations in the pore water at its edge never exceeded  $0.5 \text{ mg L}^{-1}$  between July 2002 and July 2003 (Lemke, 2004). Considering the narrow C/N ratio of surface peat at the center and the edge of the Pietzmoor and the low degree of decomposition at the edge (Table 1), it is possible that the peat itself provided the N source for denitrification.

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Schiller and Hastie (1996) report N<sub>2</sub>O release from the destruction of surface moss following clearfelling, so it is possible that the moss is the N source. This is in line with the findings by Lamers et al. (2000), who found that, at an atmospheric N deposition rate of 12 to 18 kg ha<sup>-1</sup> yr<sup>-1</sup>, excess N is accumulated in *Sphagnum* tissue, stored as free N or N-rich free amino acids. Our C/N ratio of 30 is not far from the threshold C/N ratio of 25 for significant N<sub>2</sub>O emissions reported by Klemetsson et al. (2005). In Canadian bogs and the Pietzmoor, Glatzel et al. (2004, 2006) found high CO<sub>2</sub> release rates from poorly decomposed surface *Sphagnum* peat. Since CO<sub>2</sub> release involves N mobilization and moderately dry conditions are accompanied by strong CO<sub>2</sub> emissions (Glatzel et al., 2006), in phases of moderate dryness, NO<sub>3</sub> could be accumulated that is subject to denitrification and N<sub>2</sub>O release during subsequent wetter phases.

The decreasing N<sub>2</sub>O release from the collar with the large *Eriophorum vaginatum* tussock with the progressing season suggests a competition for excess nitrogen (Silvan et al. 2005). By the end of August, plant uptake of NO<sub>3</sub> keeps N<sub>2</sub>O emission close to 0. This mechanism has been noted by Glatzel and Stahr (2001), where it led to soil N<sub>2</sub>O uptake. It is interesting that this pattern occurred only where the collar was vegetated by a large cottongrass tussock and suggests effective rhizosperic N uptake. It is likely that the wet summer favored rapid plant uptake of NO<sub>3</sub> as high soil moisture was found to be connected to efficient N uptake of *Phalaris arundinacea* (Rückauf et al., 2004).

## 4.2 CH<sub>4</sub> and N<sub>2</sub>O release from mesocosms

Gas fluxes from mesocosms were higher than from the field. This is due to constantly warm temperatures in the laboratory (Regina et al., 1999) and could, despite the two month equilibration phase, also be a consequence of an enduring disturbance effect following field sampling. As disturbance effects are site specific and there is no standard equilibration period, the comparison of the absolute magnitude of gas fluxes from mesocosms is not useful. Thus, the purpose of CH<sub>4</sub> and N<sub>2</sub>O flux determinations from mesocosms is the evaluation of differences between our treatments.

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#### 4.2.1 Methane

The variability of CH<sub>4</sub> fluxes from all mesocosms before drainage and from the control was high, but as a consequence, CH<sub>4</sub> release from the control mesocosms was not different from the mesocosms that were to be manipulated. Our finding that a water table drawdown brings CH<sub>4</sub> release to an end confirms the conclusion of the Jungkunst and Fiedler (2007) review that CH<sub>4</sub> flux rates at a water table below –10 cm are negligible (in terms of global warming potential). Strack and Waddington (2007) report a more differentiated CH<sub>4</sub> release pattern as a result of water table drawdown. They show that CH<sub>4</sub> release from hummocks may rise following a drawdown due to peat subsidence. CH<sub>4</sub> release following drainage to –50 cm also did not decline to zero (Moore and Dalva, 1993), but the peat columns sampled by Moore and Dalva were 80 cm in length. Our experimental design however eliminated the anaerobic zone, although anaerobic pockets may have been preserved, so differences due to a differing capacity for CH<sub>4</sub> oxidation one might have been able to find in the bog could not be detected. It is still interesting that immediately following the beginning of drainage, CH<sub>4</sub> fluxes at all mesocosms declined to close to 0. Also, CH<sub>4</sub> release did not reappear during the third phase. This confirms findings by Freeman et al. (2002) who reports a suppression of CH<sub>4</sub> for >1 month following a drought and Segers (1998) stated that, due slow growth rates, methanogens require a long regeneration period following exposition to oxygen. So we are not able to report a hysteresis in CH<sub>4</sub> release for the falling and rising limb as detected by Moore and Dalva (1993).

#### 4.2.2 Nitrous oxide

N<sub>2</sub>O fluxes from mesocosms declined with drainage, but did not fully recover following drainage. Increasing N<sub>2</sub>O release following drainage has been observed in field and laboratory experiments (Freeman et al., 1992; Martikainen et al., 1993; Regina et al., 1999). Dowrick et al. (1999) found that a moderate drought (with a water table at –8 cm) did not affect N<sub>2</sub>O released compared to waterlogging and that a more extreme

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drought (like the one that we simulated) causes an exponential increase in  $\text{N}_2\text{O}$  release with water table depth. On the other hand, Nykänen et al. (2002) determined very low  $\text{N}_2\text{O}$  release rates although the water table subsided up to  $-40\text{ cm}$  and one site had been fertilized with  $100\text{ kg N ha}^{-1}$  prior to the experiment. Nykänen et al. (2002) explain the low  $\text{N}_2\text{O}$  emission despite fertilization with plant uptake and the accumulation of ammonium ( $\text{NH}_4$ ) below the root zone. Another reason for this is probably the low background N load of  $6\text{ kg ha}^{-1}\text{ yr}^{-1}$  and some capacity of the peat for adsorption of  $\text{NH}_4$ . This is a profound difference to N dynamics of boreal bogs compared to temperate bogs in industrialized regions with high atmospheric N deposition and N loaded peat (Lamers et al., 2000).

There is a (non significant) rise of  $\text{N}_2\text{O}$  emissions from the manipulated mesocosm in the post-drainage phase. This could be a consequence of nitrification and an accumulation of  $\text{NO}_3$  during the drainage phase and denitrification as the water table rises again, explaining the high  $\text{NO}_3$  concentration in the pore water of the Pietzmoor during the drought in 2003 (Glatzel et al., 2006). Updegraff et al. (1995) emphasized the relationship between drainage and N mineralization. Regina et al. (1999) elaborate the link between drainage, high  $\text{NO}_3$  accumulation and increased  $\text{N}_2\text{O}$  release as well as lower  $\text{NO}_3$  concentrations and  $\text{N}_2\text{O}$  release as a consequence of rewetting. Van Beek et al. (2004) concluded that in low-land areas, ground water levels tend to control the magnitude of N losses via denitrification. In summary, although we do not know the reason for the rise of  $\text{N}_2\text{O}$  emissions in the third phase, there is evidence for denitrification following  $\text{NO}_3$  accumulation.

#### 4.3 $\text{CO}_2$ evolution from incubated disturbed samples

The purpose of laboratory incubations is the isolation of confounding factors and the absolute values obtained by this type of experiment do not approximate field fluxes. Still, Moore and Dalva (1997) suggested that integrated potential production rates and field fluxes might be similar. In any case,  $\text{CO}_2$  productions rates from peats do not differ strongly and can be compared (Glatzel et al., 2004).

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The large variability of CO<sub>2</sub> release within the unfertilized and fertilized peat is due to the wide range of peat samples used for the experiment, involving poorly as well as strongly decomposed peat as well as hummock and hollow peat. The absence of any N limitation at optimal peat moisture shows that there is no N limitation of decomposition.

5 Thus, the high N deposition rates in the region do not necessarily directly enhance peat decay, but favor N accumulation in the bog (Lamers et al., 2000). Besides the consequences on CH<sub>4</sub> and N<sub>2</sub>O release discussed above, a change in species composition is to be expected in case of persistent high N deposition and drought stress. Specifically, the competitiveness of *Sphagnum* spp. (Lamers et al., 2000; Limpens et al., 2003; Tomassen et al., 2003), *Calluna vulgaris* (Heil and Bruggink, 1987), and *Erica tetralix* (Aerts and Berendse, 1988) suffers facing atmospheric N deposition and N mineralization due to water table subsidence in favor of *Molinia caerulea* (Lamers et al., 2000; Limpens et al., 2003; Tomassen et al., 2003; Heil and Bruggink, 1987; Aerts and Berendse, 1988) and *Betula pubescens* (Tomassen et al., 2003).

## 15 5 Conclusions

Our investigations contribute to the understanding of C and N biogeochemistry in N loaded restoring peat bogs. Specifically, we were able to clarify some effects of environmental stress factors on GHG release. We captured the sensitivity of CH<sub>4</sub> and N<sub>2</sub>O fluxes to water table manipulations. In addition to the well-known water table control on CH<sub>4</sub> release, we contribute to the evidence of the water table control of N<sub>2</sub>O emissions. However, our examinations show that this control is modified by additional factors. Thus, the first part of our first hypothesis – drought decreases the CH<sub>4</sub> and N<sub>2</sub>O release – is accepted. We were not conclusively able to accept the second part of the first hypothesis – rewetting temporarily increases CH<sub>4</sub> and N<sub>2</sub>O release.

25 One of the additional factors that modify the response of CH<sub>4</sub> and N<sub>2</sub>O fluxes to water table is the degree of decomposition. We add additional evidence to the notion that the von Post humification index can be used to explain CH<sub>4</sub> and N<sub>2</sub>O release

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from restoring bogs. Here, a large variation of the humification index occurs within small areas. In the strongly decomposed center with scarce *Sphagnum* coverage, N<sub>2</sub>O release is lower than at the poorly decomposed edge with fresh N-rich *Sphagnum*. Thus, the second hypothesis, – decomposition controls CH<sub>4</sub> and N<sub>2</sub>O release – can be accepted regardless of the von Post index: if stable peat carbon compounds are available, CH<sub>4</sub> production is lower compared to easily degradable compounds which can be provided by root exudates or fresh litter. N<sub>2</sub>O emission could be enhanced when N-rich plant tissue is available for decomposition.

Our work also examined the effects of N addition to surface peat and leads to the rejection of the third hypothesis – atmospheric N deposition accelerates the decomposition of surface peat.

The ongoing restoration process in the Pietzmoor aims at the restoration of peatland ecosystems including reestablishment of natural vegetation cover, especially *Sphagnum* mosses, and of the hydrological regime (Rocheftort and Lode, 2001). Finally, the return of its functions e.g. accumulation of carbon and nutrient cycling is aspired. Realistically, this is only possible when aiming at developing an eutrophic ecosystem rather than restoring an oligotrophic one.

In summary, the examinations presented here extend our knowledge on the links between environmental stress, decomposition, methane and nitrous oxide dynamics and vegetation. On the one hand, environmental stress factors (water table and atmospheric N deposition) control GHG release, decomposition and vegetation composition. On the other hand, vegetation and decomposition control GHG release. They are part of a complex feedback loop that drives the dynamics of GHG emissions in N saturated restoring bogs. Concerning GHG exchange, future research in N saturated restoring bogs needs to compare two phenomena: elevated GHG release from poorly decomposed fresh peat and elevated methane release due to a rising proportion of sedges on strongly decomposed peat.

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**Table 1.** Carbon (C) and nitrogen (N) concentration and von Post humification index in the top 15 cm of peat cores used for the water table manipulation experiment from the Pietzmoor, Germany. Mean values and standard deviation from six replicates are shown. Significantly different values (Wilcoxon test) within one line are marked with different letters.

	Center	Edge %
C (%)	48.28±0.69 a	44.10±0.60 b
N (%)	1.59±0.09 a	1.39±0.11 b
	dimensionless	
C/N ratio	30.42±1.54 a	31.92±2.42 a
von Post index	5.3±0.8 a	2.7±0.7 b

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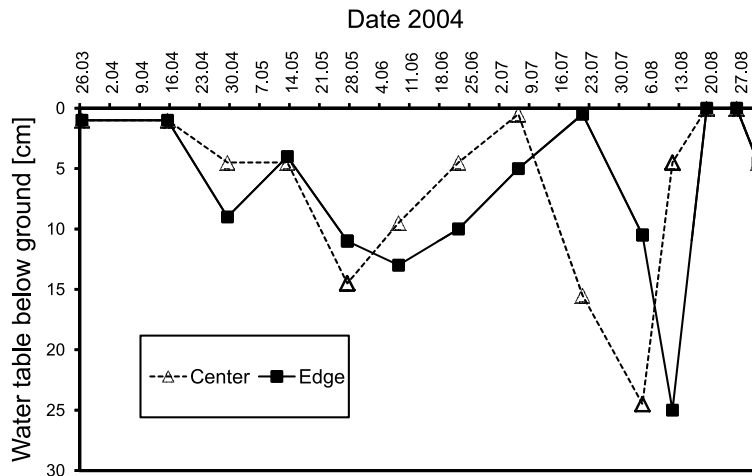
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**Fig. 1.** Water table at the center and at the edge of the research site in the Pietzmoor from March to August 2004.

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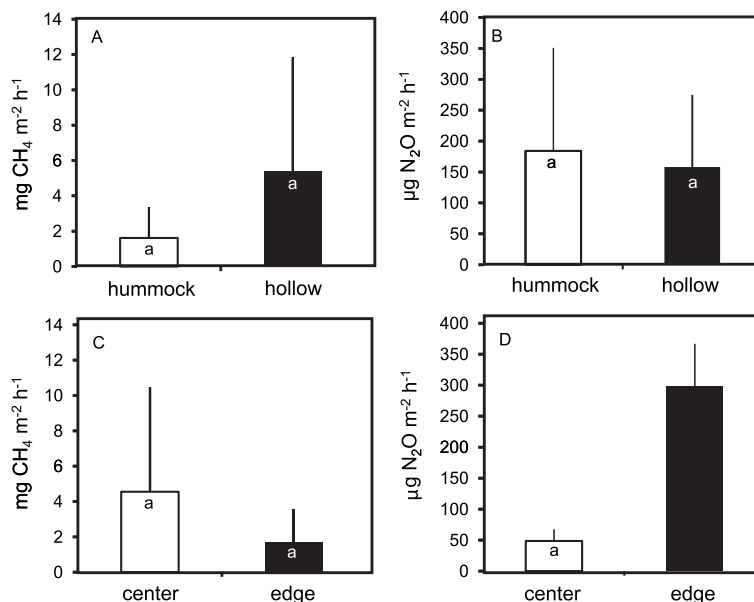
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**Fig. 2.** Methane (CH<sub>4</sub>) and nitrous oxide (N<sub>2</sub>O) release from hummocks and hollows at the center and the edge of the Pietzmoor, Germany. Shown are means and standard deviations of CH<sub>4</sub> and N<sub>2</sub>O release of 14 field gas flux determinations from March to August 2004 from six hummocks and four hollows (CH<sub>4</sub>: A, N<sub>2</sub>O: B) originating from five center and five edge locations (CH<sub>4</sub>: C, N<sub>2</sub>O: D). Significantly different values (Wilcoxon test) are marked with different letters.

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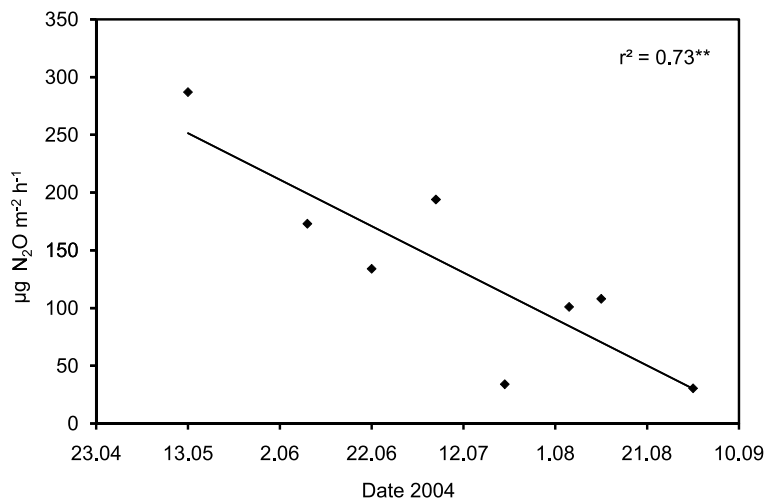
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**Fig. 3.** Nitrous oxide (N<sub>2</sub>O) release from an *Eriophorum vaginatum* dominated hummock from April to August 2004 and its relation to sampling date at the edge of the Pietzmoor, Germany.

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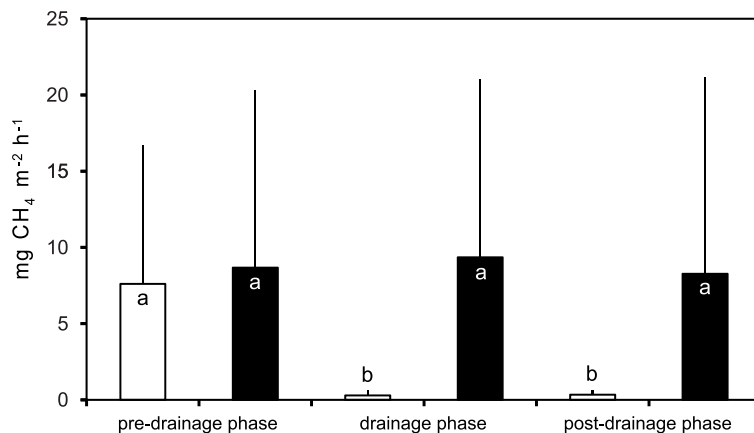
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**Fig. 4.** Methane ( $\text{CH}_4$ ) release from peat cores before, during, and following the drainage experiment (open bars). Unmanipulated control cores are black. Mean values and standard deviation from three replicates are shown. Significantly different values (Wilcoxon test) are marked with different letters.

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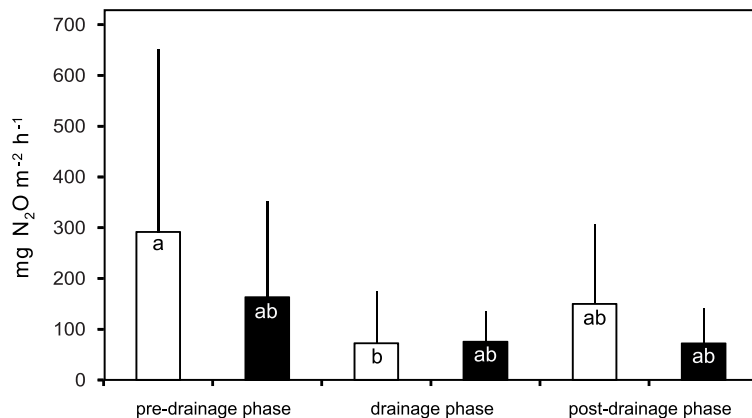
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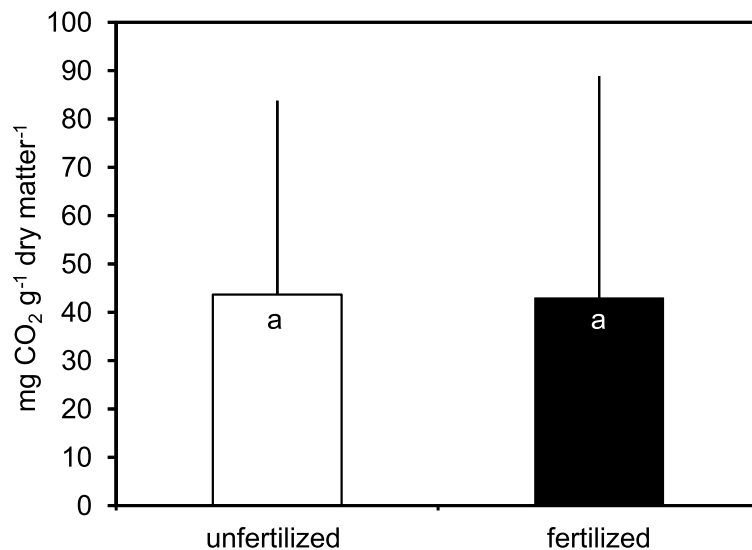
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**Fig. 5.** Nitrous oxide (N<sub>2</sub>O) release from peat cores before, during, and following the drainage experiment (open bars). Unmanipulated control cores are black. Mean values and standard deviation from three replicates are shown. Significantly different values (Wilcoxon test) are marked with different letters.

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**Fig. 6.** Carbon dioxide (CO<sub>2</sub>) release from unfertilized peat and peat fertilized with ammonium nitrate from the Pietzmoor, Germany, following a 42 day incubation. Mean values and standard deviation from 17 samples are shown. Significantly different values (Wilcoxon test) are marked with different letters.

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